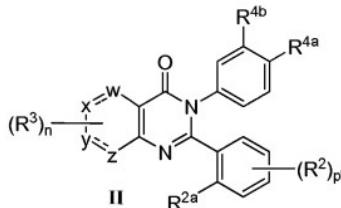


In the claims:

1. Cancelled

2. (Currently amended) The compound according to Claim 1 of the formula II:



wherein  $a$ ,  $w$ ,  $x$ ,  $y$ ,  $z$ , dashed line,  $R^3$ ,  $R^4$ ,  $R^6$  and  $R^7$  are defined as in Claim 1 for the compound of the Formula I; and

w, x, y and z are independently selected from CH or CH<sub>2</sub>;

a dashed line represents an optional double bond:

a is 0 or 1;

b is 0 or 1;

m is 0, 1, or 2;

n is 0 to 2;

r is 0 or 1;

s is 0 or 1;

n is 0 or 1;

p' is 0 to 2;

R<sup>2</sup> is selected from:

- 1)       $(C=O)_a C_1-C_{10}$  alkyl,
- 2)       $(C=O)_a aryl,$
- 3)       $(C=O)_a NR_6R^7,$
- 4)       $(C=O)_a C_3-C_8$  cycloalkyl,
- 5)       $(C=O)_a heterocyclyl,$

- 6)  $\text{SO}_2\text{NR}^6\text{R}^7$ , and  
 7)  $\text{SO}_2\text{C}_1\text{-C}_{10}$  alkyl,

said alkyl, aryl, cycloalkyl, and heterocyclyl is optionally substituted with one or more substituents selected from R<sup>4</sup>;

R<sup>2a</sup> is selected from: halogen and (C<sub>1</sub>-C<sub>6</sub>)alkyl; and

R<sup>3</sup> is selected from:

- 1)  $(\text{C}=\text{O})_a\text{ObC}_1\text{-C}_{10}$  alkyl,
- 2)  $(\text{C}=\text{O})_a\text{Obaryl}$ ,
- 3)  $(\text{C}=\text{O})_a\text{ObC}_2\text{-C}_{10}$  alkenyl,
- 4)  $(\text{C}=\text{O})_a\text{ObC}_2\text{-C}_{10}$  alkynyl,
- 5)  $\text{CO}_2\text{H}$ ,
- 6) halo,
- 7) OH,
- 8)  $\text{ObC}_1\text{-C}_6$  perfluoroalkyl,
- 9)  $(\text{C}=\text{O})_a\text{NR}^6\text{R}^7$ ,
- 10) CN,
- 11)  $(\text{C}=\text{O})_a\text{ObC}_3\text{-C}_8$  cycloalkyl,
- 12)  $(\text{C}=\text{O})_a\text{Obheterocyclyl}$ ,
- 13)  $\text{SO}_2\text{NR}^6\text{R}^7$ , and
- 14)  $\text{SO}_2\text{C}_1\text{-C}_{10}$  alkyl,

said alkyl, aryl, alkenyl, alkynyl, cycloalkyl, and heterocyclyl is optionally substituted with one or more substituents selected from R<sup>4</sup>;

R<sup>4</sup> is selected from:

- 1)  $(\text{C}=\text{O})_a\text{ObC}_1\text{-C}_{10}$  alkyl,
- 2)  $(\text{C}=\text{O})_a\text{Obaryl}$ ,
- 3)  $\text{C}_2\text{-C}_{10}$  alkenyl,
- 4)  $\text{C}_2\text{-C}_{10}$  alkynyl,
- 5)  $(\text{C}=\text{O})_a\text{Ob}$  heterocyclyl,
- 6)  $\text{CO}_2\text{H}$ ,
- 7) halo,
- 8) CN,
- 9) OH,
- 10)  $\text{ObC}_1\text{-C}_6$  perfluoroalkyl,

- 11)      O<sub>a</sub>(C=O)<sub>b</sub>NR<sub>6</sub>R<sub>7</sub>,
- 12)      oxo,
- 13)      CHO,
- 14)      (N=O)R<sub>6</sub>R<sub>7</sub>, or
- 15)      (C=O)<sub>a</sub>O<sub>b</sub>C<sub>3</sub>-C<sub>8</sub> cycloalkyl,
- 16)      SO<sub>2</sub>C<sub>1</sub>-C<sub>10</sub>alkyl,
- 17)      SO<sub>2</sub>NR<sub>6</sub>R<sub>7</sub>,

said alkyl, aryl, alkenyl, alkynyl, heterocycl<sub>l</sub>, and cycloalkyl optionally substituted with one or more substituents selected from R5:

R<sup>4a</sup> and R<sup>4b</sup> are independently selected from: hydrogen, halogen and (C<sub>1</sub>-C<sub>6</sub>)alkyl, provided that at least one is not hydrogen, or

R<sup>4a</sup> and R<sup>4b</sup> are combined to form a diradical selected from -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-,  
-CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-,-CH=CH-O- and -CH=CH-N-

R5 is selected from:

- 1)      (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>1</sub>-C<sub>10</sub>)alkyl,
- 2)      O<sub>f</sub>(C<sub>1</sub>-C<sub>3</sub>)perfluoroalkyl,
- 3)      (C<sub>0</sub>-C<sub>6</sub>)alkylene-S(O)<sub>m</sub>R<sup>a</sup>,
- 4)      oxo,
- 5)      OH,
- 6)      halo,
- 7)      CN,
- 8)      (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>2</sub>-C<sub>10</sub>)alkenyl,
- 9)      (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>2</sub>-C<sub>10</sub>)alkynyl,
- 10)     (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>3</sub>-C<sub>6</sub>)cycloalkyl,
- 11)     (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>0</sub>-C<sub>6</sub>)alkylene-aryl,
- 12)     (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>0</sub>-C<sub>6</sub>)alkylene-heterocycl<sub>l</sub>,
- 13)     (C=O)<sub>1</sub>O<sub>S</sub>(C<sub>0</sub>-C<sub>6</sub>)alkylene-N(R<sup>b</sup>)<sub>2</sub>,
- 14)     C(O)R<sup>a</sup>,
- 15)     (C<sub>0</sub>-C<sub>6</sub>)alkylene-CO<sub>2</sub>R<sup>a</sup>,
- 16)     C(O)H,
- 17)     (C<sub>0</sub>-C<sub>6</sub>)alkylene-CO<sub>2</sub>H, and
- 18)     C(O)N(R<sup>b</sup>)<sub>2</sub>,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocycl is optionally substituted with up to three substituents selected from R<sup>b</sup>, OH, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, halogen, CO<sub>2</sub>H, CN, O(C=O)C<sub>1</sub>-C<sub>6</sub> alkyl, oxo, and N(R<sup>b</sup>)<sub>2</sub>:

R<sup>6</sup> and R<sup>7</sup> are independently selected from:

- 1) H,
- 2) (C=O)O<sub>b</sub>C<sub>1</sub>-C<sub>10</sub> alkyl,
- 3) (C=O)O<sub>b</sub>C<sub>3</sub>-C<sub>8</sub> cycloalkyl,
- 4) (C=O)O<sub>b</sub>aryl,
- 5) (C=O)O<sub>b</sub>heterocyclyl,
- 6) C<sub>1</sub>-C<sub>10</sub> alkyl,
- 7) aryl,
- 8) C<sub>2</sub>-C<sub>10</sub> alkenyl,
- 9) C<sub>2</sub>-C<sub>10</sub> alkynyl,
- 10) heterocyclyl,
- 11) C<sub>3</sub>-C<sub>8</sub> cycloalkyl,
- 12) SO<sub>2</sub>R<sup>a</sup>, and
- 13) (C=O)NR<sup>b</sup>,

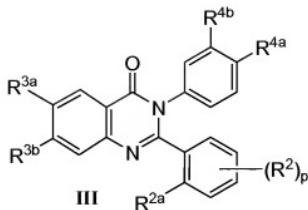
said alkyl, cycloalkyl, aryl, heterocycl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R<sup>5</sup>, or

R<sup>6</sup> and R<sup>7</sup> can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R<sup>5</sup>

R<sup>a</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, aryl, or heterocycl; and

R<sup>b</sup> is H, (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>1</sub>-C<sub>6</sub>)alkyl-NR<sup>a</sup><sub>2</sub>, (C<sub>1</sub>-C<sub>6</sub>)alkyl-NH<sub>2</sub>, (C<sub>1</sub>-C<sub>6</sub>)alkyl-NHR<sup>a</sup>, aryl, heterocyclyl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C=O)OC<sub>1</sub>-C<sub>6</sub> alkyl, (C=O)C<sub>1</sub>-C<sub>6</sub> alkyl or S(O)<sub>2</sub>R<sup>a</sup>.

3. (Original) A compound of the formula III, or a pharmaceutically acceptable salt or stereoisomer thereof,



wherein

b is 0 or 1;

m is 0, 1 or 2;

p' is 0 to 2;

r is 0 or 1;

s is 0 or 1;

R<sup>2</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkylene-NR<sub>6</sub>R<sup>7</sup>; said alkylene is optionally substituted with up to three substituents selected from OH, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, halogen, CO<sub>2</sub>H, CN, O(C=O)C<sub>1</sub>-C<sub>6</sub> alkyl, oxo, and NR<sub>6</sub>R<sup>7</sup>;

R<sup>2a</sup> is selected from: halogen and (C<sub>1</sub>-C<sub>6</sub>)alkyl;

R<sup>3a</sup> and R<sup>3b</sup> are independently selected from: hydrogen and halogen; and

R<sup>4a</sup> and R<sup>4b</sup> are independently selected from: hydrogen, halogen, and (C<sub>1</sub>-C<sub>6</sub>)alkyl, provided that at least one is not hydrogen;

R<sup>5</sup> is selected from:

- 1) (C=O)<sub>r</sub>O<sub>S</sub>(C<sub>1</sub>-C<sub>10</sub>)alkyl,
- 2) O<sub>r</sub>(C<sub>1</sub>-C<sub>3</sub>)perfluoroalkyl,
- 3) (C<sub>0</sub>-C<sub>6</sub>)alkylene-S(O)<sub>m</sub>R<sup>a</sup>,
- 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) (C=O)<sub>r</sub>O<sub>S</sub>(C<sub>2</sub>-C<sub>10</sub>)alkenyl,

- 9)  $(C=O)_tOs(C_2-C_{10})alkynyl,$
- 10)  $(C=O)_tOs(C_3-C_6)cycloalkyl,$
- 11)  $(C=O)_tOs(C_0-C_6)alkylene-aryl,$
- 12)  $(C=O)_tOs(C_0-C_6)alkylene-heterocyclyl,$
- 13)  $(C=O)_tOs(C_0-C_6)alkylene-N(R^b)_2,$
- 14)  $C(O)R^a,$
- 15)  $(C_0-C_6)alkylene-CO_2R^a,$
- 16)  $C(O)H,$
- 17)  $(C_0-C_6)alkylene-CO_2H,$  and
- 18)  $C(O)N(R^b)_2,$

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R<sup>b</sup>, OH, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, halogen, CO<sub>2</sub>H, CN, O(C=O)C<sub>1</sub>-C<sub>6</sub> alkyl, oxo, and N(R<sup>b</sup>)<sub>2</sub>;

R<sup>6</sup> and R<sup>7</sup> are independently selected from:

- 1) H,
- 2)  $(C=O)ObC_1-C_{10}$  alkyl,
- 3)  $(C=O)ObC_3-C_8$  cycloalkyl,
- 4)  $(C=O)Obaryl,$
- 5)  $(C=O)Obheterocyclyl,$
- 6) C<sub>1</sub>-C<sub>10</sub> alkyl,
- 7) aryl,
- 8) C<sub>2</sub>-C<sub>10</sub> alkenyl,
- 9) C<sub>2</sub>-C<sub>10</sub> alkynyl,
- 10) heterocyclyl,
- 11) C<sub>3</sub>-C<sub>8</sub> cycloalkyl,
- 12) SO<sub>2</sub>R<sup>a</sup>, and
- 13)  $(C=O)NR^b_2,$

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R<sup>5</sup>, or

R<sup>6</sup> and R<sup>7</sup> can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R<sup>5</sup>;

R<sup>a</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, aryl, or heterocycl; and

R<sup>b</sup> is H, (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>1</sub>-C<sub>6</sub>)alkyl-NR<sup>a</sup><sub>2</sub>, (C<sub>1</sub>-C<sub>6</sub>)alkyl-NH<sub>2</sub>, (C<sub>1</sub>-C<sub>6</sub>)alkyl-NHRA<sup>a</sup>, aryl, heterocycl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C=O)OC<sub>1</sub>-C<sub>6</sub> alkyl, (C=O)C<sub>1</sub>-C<sub>6</sub> alkyl or S(O)<sub>2</sub>R<sup>a</sup>.

4. (Original) The compound according to Claim 3, or the pharmaceutically acceptable salt or stereoisomer thereof, wherein p', R<sup>2a</sup>, R<sup>3a</sup>, R<sup>3b</sup>, R<sup>4a</sup>, R<sup>4b</sup> and R<sup>5</sup> are as defined for Formula III in Claim 3 and

R<sup>2</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkylene-NR<sup>6</sup>R<sup>7</sup>;

R<sup>6</sup> and R<sup>7</sup> are independently selected from:

- 1) H,
- 2) C<sub>1</sub>-C<sub>10</sub> alkyl,
- 3) aryl,
- 4) heterocycl,
- 5) C<sub>2</sub>-C<sub>10</sub> alkenyl,
- 6) C<sub>2</sub>-C<sub>10</sub> alkynyl, and
- 7) C<sub>3</sub>-C<sub>8</sub> cycloalkyl,

said alkyl, cycloalkyl, aryl, heterocycl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R<sup>5</sup>, or

R<sup>6</sup> and R<sup>7</sup> can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R<sup>5</sup>.

5. (Original) A compound selected from:

2-(2-bromophenyl)-3-(4-methylphenyl)quinazolin-4(3H)-one;

2-(2-bromophenyl)-3-(4-methylphenyl)-quinazolin-4(3H)-one;

2-(2-chlorophenyl)-3-(4-methylphenyl)-quinazolin-4(3H)-one;

2-(2,4-dichlorophenyl)-3-(4-methylphenyl)quinazolin-4(3H)-one;

2-(2-bromophenyl)-3-(4-chlorophenyl)-quinazolin-4(3H)-one;  
2-(2-bromophenyl)-3-(3-fluoro-4-methylphenyl)-quinazolin-4(3H)-one;  
3-(3a,7a-dihydro-1H-indol-5-yl)-2-(2-bromophenyl)-quinazolin-4(3H)-one;  
6-chloro-2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)-quinazolin-4(3H)-one;  
2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;  
2-(2-methylphenyl)-3-(4-methylphenyl)-quinazolin-4(3H)-one;  
7-chloro-2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;  
2-(2-bromophenyl)-7-chloro-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;  
7-chloro-2-(2-chlorophenyl)-3-(1H-indol-5-yl)quinazolin-4(3H)-one;  
2-(2-bromophenyl)-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;  
2-(2-bromophenyl)-3-(3-fluoro-4-methyl-phenyl)pyrido[2,3-d]pyrimidin-4(3H)-one;  
2-(5-bromo-2-chlorophenyl)-7-chloro-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;  
2-(4-bromo-2-chlorophenyl)-7-chloro-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;  
2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)-5,6,7,8-tetrahydroquinazolin-4(3H)-one;  
7-chloro-2-{2-chloro-3-[(dimethylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one ;  
7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(4-methylpiperazin-1-yl)methyl]phenyl}quinazolin-4(3H)-one;  
7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(methylamino)methyl]-phenyl}quinazolin-4(3H)-one;  
7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(4-methylpiperazin-1-yl)methyl]phenyl}quinazolin-4(3H)-one;  
7-chloro-2-{2-chloro-3-[(ethylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;  
7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(isopropylamino)methyl]-phenyl}quinazolin-4(3H)-one;

7-chloro-2-{2-chloro-3-[(cyclobutylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

2-[3-(azetidin-1-ylmethyl)-2-chlorophenyl]-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(pyrrolidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-[(3S)-3-hydroxypyrrolidin-1-yl]methyl)phenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-[(3S)-3-(methoxymethyl)pyrrolidin-1-yl]methyl)phenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(pyrrolidin-3-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(morpholin-4-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(piperidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

2-{3-[(4-aminopiperidin-1-yl)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(piperidin-4-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(4-fluoropiperidin-1-yl)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(piperazin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

2-{3-[(4-acetyl piperazin-1-yl)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-{[4-(methylsulfonyl)piperazin-1-yl]methyl}phenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-[(2-hydroxyethyl)amino]-methyl}phenyl)quinazolin-4(3H)-one;

7-chloro-2-[2-chloro-3-({[2-(dimethylamino)ethyl]amino}methyl)phenyl]-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-[(2-morpholin-4-ylethyl)amino]methyl}phenyl)quinazolin-4(3H)-one;

2-{3-[(3-aminopyrrolidin-1-yl)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-({[(1-methylpiperidin-3-yl)methyl]amino}methyl)phenyl]quinazolin-4(3H)-one;

2-(3-{{[3-(aminomethyl)-1-methyl-1lambda~5-piperidin-1-yl]methyl}-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

2-{3-[(benzylamino)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(4-methylpiperazin-1-yl)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-2-{2-chloro-5-[(ethylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(isopropylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(pyrrolidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(pyrrolidin-3-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(morpholin-4-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(piperidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(piperidin-4-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(piperazin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-5-{{4-(methylsulfonyl)piperazin-1-yl}methyl}phenyl)quinazolin-4(3H)-one; and

7-chloro-2-[2-chloro-5-({[2-(dimethylamino)ethyl]amino}methyl)phenyl]-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

or a pharmaceutically acceptable salt thereof.

6. (Currently amended) A pharmaceutical composition that is comprised of a compound in accordance with Claim 42 and a pharmaceutically acceptable carrier.

7. (Original) A pharmaceutical composition that is comprised of a compound in accordance with Claim 3 and a pharmaceutically acceptable carrier.

8. (Currently amended) A method of treating or preventing cancer in a mammal in need of such treatment that is comprised of administering to said mammal a therapeutically effective amount of a compound of Claim 42.

9. Previously cancelled

10. (Original) A method of treating cancer or preventing cancer in accordance with Claim 8 wherein the cancer is selected from cancers of the brain, genitourinary tract, lymphatic system, stomach, larynx and lung.

11. (Original) A method of treating or preventing cancer in accordance with Claim 8 wherein the cancer is selected from histiocytic lymphoma, lung adenocarcinoma, small cell lung cancers, pancreatic cancer, glioblastomas and breast carcinoma.

12.-20. Previously cancelled

21.-24. Cancelled

25.-28. Previously cancelled

29. Cancelled

30. Previously cancelled

31.-34. Cancelled